AMENDMENTS TO THE CLAIMS:

The following listing of claims replaces all prior listings, and all prior versions, of claims in the application.

LISTING OF CLAIMS:

1. (Original) A method for producing an optically active chromancarboxylate, comprising a step of esterifying a racemic chromancarboxylic acid in an organic solvent comprising an alcohol in the presence of a biocatalyst, the racemic chromancarboxylic acid being represented by the formula 1:

$$R_n = \begin{pmatrix} 0 \\ 1 \end{pmatrix} X_m$$
 (1)

wherein R is a halogen atom, a hydroxyl group, a nitro group, an amino group, a cyano group, a chloromethyl group, a trifluoromethyl group, a carboxyl group, a carboxymethyl group, a carboxyethyl group, a carboxyphenyl group, a substituted or unsubstituted alkyl group or a substituted or unsubstituted aryl group, and a plurality of R groups, if any, may be the same or different; X is a halogen atom, a hydroxyl group, a nitro group, an amino group, a cyano group, a chloromethyl group, a trifluoromethyl group, a carboxyl group, a carboxymethyl group, a carboxyethyl group, a carboxyphenyl group, a substituted or unsubstituted alkyl group or a substituted or unsubstituted aryl group with the proviso that at least one X is a carboxyl group, and a plurality of X groups, if any, may be the same or different; m is an integer of from 1 to 5; and n is an integer of from 0 to 4.

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- 2. (Original) The method according to claim 1, wherein the biocatalyst is a hydrolase produced by microorganisms.
- (Original) The method according to claim 2, wherein the hydrolase is lipase.
- 4. (Original) The method according to claim 3, wherein the lipase is derived from microorganisms belonging to genus *Candida*.
- 5. (Currently amended) The method according to <u>claimany one of claims</u> 1 to 4, wherein the alcohol has from 1 to 24 carbon atoms.
- 6. (Original) The method according to claim 5, wherein the alcohol is methanol, ethanol, n-propyl alcohol, isopropyl alcohol, n-butyl alcohol or isobutyl alcohol.
- 7. (Original) The method according to claim 6, wherein the alcohol is methanol.
- 8. (Currently amended) The method according to <u>claimany one of claims</u> 1 to 7, wherein the chromancarboxylic acid is selected from the group consisting of chroman-2-carboxylic acid, chroman-3-carboxylic acid, chroman-4-carboxylic acid, 6-hydroxychroman-2-rnethyl-2-carboxylic acid, 2-

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carboxymethyi 6-hydroxy- 2-methylchroman, 6-hydroxy- 5-methylchroman-2-carboxylic acid, 6-hydroxy-7,8-dimethylchroman-2-carboxylic acid, 6-hydroxy-2,7,8-trimethylchroman-2-carboxylic acid, 6-hydroxy-2, 7, 8-trimethylchroman-2-ylpropionic acid and 6-hydroxy-2, 5,7,8-tetramethylchroman-2-carboxylic acid.

- 9. (Original) The method according to claim 8, wherein the chromancarboxylic acid is selected from the group consisting of chroman-2-carboxylic acid, 6 hydroxy-2, 7,8-trimethyl-2-carboxymethylchroman, 6-hydroxy-2, 7,8-trimethylchroman- 2-ylpropionic acid and 6-hydroxy-2, 5,7,8-tetramethylchroman-2-carboxylic acid.
- 10. (Original) The method according to claim 9, wherein the chromancarboxylic acid is 6-hydroxy-2,5,7,8 -tetramethylchroman-2-carboxylic acid.
- 11. (Currently amended) The method according to <u>claimany one of claims</u> 1 to 10, further comprising a step of separating a mirror image of the chromancarboxylic acid which is converted into the optically active chromancarboxylate from a reaction production solution after the esterification.
- 12. (Currently amended) The method according to <u>claimany one of claims</u> 1 to 11, further comprising a step of hydrolyzing the optically active chromancarboxylate.